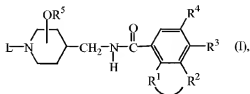


This listing of claims will replace all prior versions, and listings, of claims in the application.

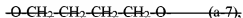
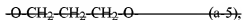
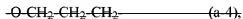
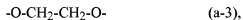
Listing of Claims:

1. (Currently Amended) A compound of formula (I)



a stereochemically isomeric form thereof, an *N*-oxide form thereof, or a pharmaceutically acceptable acid or base addition salt thereof, wherein

-R¹-R²- is a bivalent radical of formula



wherein in said bivalent radicals optionally one or two hydrogen atoms on the same or a different carbon atom may be replaced by C₁₋₆alkyl or hydroxy,

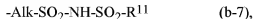
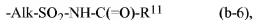
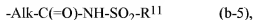
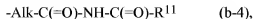
R³ is hydrogen, halo, C₁₋₆alkyl or C₁₋₆alkyloxy;

R⁴ is hydrogen, halo, C₁₋₆alkyl; C₁₋₆alkyl substituted with cyano, or C₁₋₆alkyloxy;


C₁₋₆alkyloxy; cyano; amino or mono or di(C₁₋₆alkyl)amino;

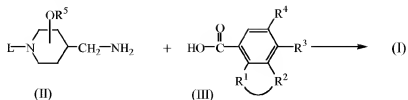
R⁵ is hydrogen or C₁₋₆alkyl, and the -OR⁵ radical is situated at the 3- or 4-position of the piperidine moiety;

L is a radical of formula

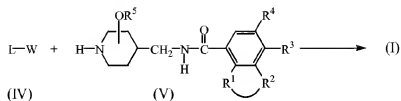


wherein each Alk is C₁₋₁₂alkanediyl; and
R⁶ is aminosulfonyl optionally substituted with C₁₋₄alkyl, C₃₋₆cycloalkyl or phenyl;
R⁷ is C₁₋₆alkylsulfonyl;
X is NR⁸; said R⁸ being C₁₋₆alkyl;
R⁹ is C₁₋₆alkylsulfonylamino;
Y is a O, S, or NR¹⁰ wherein R¹⁰ is hydrogen or C₁₋₆alkyl; and
R¹¹ is C₁₋₆alkyl or phenyl wherein the -OR⁵ radical is situated at the 3-position.

-  2. (Previously Presented) The compound as claimed in claim 1 wherein the -OR⁵ radical is situated at the 3-position of the piperidine moiety ^{has} having the trans configuration.
3. (Previously Presented) The compound as claimed in claim 2 wherein the absolute configuration of said piperidine moiety is (3S, 4S).
4. (Previously Presented) The compound as claimed in claim 1 wherein L is a radical of formula (b-1) wherein Alk is C₁₋₄alkanediyl, and R⁶ aminosulfonyl or aminosulfonyl substituted with C₁₋₄alkyl or phenyl.
5. (Previously Presented) The compound as claimed in claim 1 wherein L is a radical (b-5) wherein Alk is C₁₋₄alkanediyl, and R¹¹ is C₁₋₄alkyl.
6. (Previously Presented) The compound as claimed in claim 1 wherein L is a radical (b-7) wherein Alk is C₁₋₄alkanediyl, and R¹¹ is C₁₋₄alkyl.
7. (Previously Presented) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a therapeutically active amount of a compound according to claim 1.
8. (Canceled)
9. (Canceled)
- ⁸
10. (Original) A process for preparing a compound of formula (I) wherein
a) an intermediate of formula (II) is reacted with an carboxylic acid derivative of formula (III) or a reactive functional derivative thereof;



b) an intermediate of formula (IV) is *N*-alkylated with an intermediate of formula (V), in a reaction-inert solvent and, optionally in the presence of a suitable base;



wherein in the above reaction schemes the radicals -R¹-R²-, R³, R⁴, R⁵, and L are as defined in claim 1 and W is an appropriate leaving group;

c) or, compounds of formula (I) are converted into each other following art-known transformation reactions; or if desired; a compound of formula (I) is converted into a pharmaceutically acceptable acid addition salt, or conversely, an acid addition salt of a compound of formula (I) is converted into a free base form with alkali; and, if desired, preparing stereochemically isomeric forms thereof.

11. (Withdrawn) A method for the treatment of ~~5HT₂-related~~ disorders comprising administering to a patient ~~in need thereof~~ an effective amount of a compound according to claim 1.

9
12. (Withdrawn) A method for treating patients suffering from gastrointestinal ~~conditions~~ disorders comprising administering to the patient an effective amount of a compound according to claim 1.

10
13. (Withdrawn) A method for treating hypermotility, irritable bowel syndrome, constipation or diarrhea predominant IBS, pain and non-pain predominant IBS and bowel

DOCKET NO.: JANM-0773/PRD2061USPT

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Application No.: 10/560,300

Office Action Dated: June 9, 2009

hypersensitivity comprising administering to a patient in need thereof an effective amount of a compound according to claim 1.